To: Rusyn, Ivan[IRusyn@cvm.tamu.edu]; Kate Guyton[GuytonK@iarc.fr]; Martin, Matt[Martin.Matt@epa.gov]; dmreif@ncsu.edu[dmreif@ncsu.edu]
From: Chiu, Weihsueh
Sent: Mon 2/27/2017 1:53:40 AM
Subject: Fwd: RTP-16-407R2: Final Decision

Bad news...

Regards,
Weihsueh
==========
Weihsueh A. Chiu, PhD
Professor
Texas A&M University

-- Sent from my mobile device. Please excuse terse responses.

Begin forwarded message:

From: "RTP (ELS)" <eesserver@eesmail.elsevier.com>
Date: February 26, 2017 at 7:36:49 PM CST
To: <wchiu@cvm.tamu.edu>
Subject: RTP-16-407R2: Final Decision
Reply-To: "RTP (ELS)" <rtp@elsevier.com>

Ms. No.: RTP-16-407R2
Title: Use of high-throughput in vitro toxicity screening data in cancer hazard evaluations by IARC Monograph Working Groups
Corresponding Author: Dr. Weihsueh A Chiu
Authors: Kathryn Z Guyton, PhD; Matthew T Martin, PhD; David M Reif, PhD; Ivan Rusyn, MD, PhD

Dear Dr. Chiu,

Thank you for submitting your manuscript to Regulatory Toxicology and Pharmacology. Your paper, referenced above, has been reviewed by experts in the field. The reviewer comments, included below, are extensive to the point of precluding just an interim revision of your current submission, and I regret that your manuscript cannot be accepted for publication as it stands.
However, there is interest in the topic you presented and you are encouraged to resubmit a thoroughly improved manuscript, along the recommendations offered by the reviewers. In this case, the new manuscript would be considered a new submission and would be given a new manuscript number with a new date of receipt.

Again, this critique of your paper does not imply a lack of interest in this area of research, and I hope you will continue to submit your work to this journal in the future.

Sincerely,

Dr. Gio Batta Gori, Editor-in-Chief
Regulatory Toxicology and Pharmacology
E-mail: rtp@elsevier.com

Reviewers' comments: